

Company Description

VIVEbiotech is a GMP Contract Development and Manufacturing Organisation (CDMO) specialized in lentiviral vectors. Before VIVEbiotech was established, its team had already reached some key milestones within the Advanced Therapies field. This allowed VIVEbiotech to become an integral service lentiviral CDMO, and since the beginning it has focused its efforts on offering custom services to international customers.

Information

📅 Deadline: 2021-04-18
🏢 Category: Business
📍 Province: Gipuzkoa

🌐 Country: Basque Country
🏙️ City: San Sebastian

Company

VIVEBIOTECH SL



Main functions, requisites & benefits

Main functions

To design, prepare and execute development batches of lentiviral vectors. To research on process optimization. Propose changes to optimize the USP process and analyze them according the results of the development batches. To draft of protocols, standardized operating procedures (SOPs) and other relevant documents. To comply with and ensure compliance with internal work standards. To be part of a multidisciplinary team to carry out process optimizations and research. To carry out maintenance of process development zones according to internal standards.

Requisites

Background: Degree related to research in Biomedicine (Science, Biology, Biochemistry, Chemistry, Pharmacy ...) or related Engineering. Related higher degree. Ability to propose possible process improvements, propose experiments and analyze results. Previous experience in USP development of bioprocesses will be positively valued. Spoken and written communication skills (both in Spanish and English). Team working skills and ability to work under deadlines.

